Appl. No. ·

Filed

*5*9/754,949

January 4, 2001

claim 20 is that in the method of claim 20 the host cell endogenously expresses Par-4 rather than being transfected with nucleic acid encoding Par-4. As the methods of original claims 1 and 20 are very similar, they would not have a separate status in the art and the searching would be coextensive. Thus, Applicants believe that the amendment of claim 1 is appropriate in view of the restriction requirement and respectfully request examination of the application.

Respectfully submitted,

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Dated: Fuirula 3, 2002

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VERSION WITH MARKINGS TO SHOW CHANGES MADE

Claims 1 - 4, 6, 9 and 17 have been amended as follows:

1. A method for identifying inhibitors of neuronal degeneration comprising:

(aA) (1) cotransfecting eukaryotic host cells expressing a presenilin protein (PS), with a polynucleotide encoding a Par-4 polypeptide, and an NF-κB dependent reporter construct, (b2) exposing the cotransfected cells to a candidate molecule, and (e3) monitoring the ability of said candidate molecule to induce NF-κB activation; or

(B) (1) transfecting eukaryotic host cells endogenously expressing Par-4 and a presenilin (PS) protein with nucleic acid encoding an NF-κB dependent reporter construct, (2) exposing the transfected cells to a candidate molecule, and (3) monitoring the ability of said candidate molecule to induce NF-κB activation.

- 2. The method of claim 1 wherein said eukaryotic host cells of part (A) are mammalian cells endogenously expressing PS.
- 3. The method of claim 1 wherein said eukaryotic host cells of part (A) are mammalian cells transfected with nucleic acid encoding PS.
 - 4. The method of claim 31 wherein said PS is PS1.
 - 6. The method of claim 31 wherein said PS is FAD PS.
- 9. The method of claim 1 wherein said eukaryotic host cells of part (A) are neuronal cells.
- 17. The method of claim 1 wherein the <u>transfected or cotransfected cells</u> are exposed to a plurality of candidate molecules.

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